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PCT

NOTIFICATION OF ELECTION

(PCT Rule 61.2)

To:

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Date of mailing (day/month/year) 16 March 2000 (16.03.00)	
International application No. PCT/GB99/02127	**Applicant's or agent's file reference** SPG/P36002WO
International filing date (day/month/year) 02 July 1999 (02.07.99)	**Priority date** (day/month/year) 04 July 1998 (04.07.98)
Applicant PARKER, Dawood et al	

1. The designated Office is hereby notified of its election made:

[X] in the demand filed with the International Preliminary Examining Authority on:

01 February 2000 (01.02.00)

[ ] in a notice effecting later election filed with the International Bureau on:

2. The election [X] was

[ ] was not

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The International Bureau of WIPO 34, chemin des Colombettes 1211 Geneva 20, Switzerland	Authorized officer Juan Cruz
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PCT WORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 : A61B 5/00 — A1

(11) International Publication Number: WO 00/01294

(43) International Publication Date: 13 January 2000 (13.01.00)

(21) International Application Number: PCT/GB99/02127

(22) International Filing Date: 2 July 1999 (02.07.99)

(30) Priority Data:

9814464.5	4 July 1998 (04.07.98)	GB
9824899.0	13 November 1998 (13.11.98)	GB
9825243.0	19 November 1998 (19.11.98)	GB

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(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published

With international search report.

(54) Title: NON-INVASIVE MEASUREMENT OF BLOOD ANALYTES

(57) Abstract

There is described a device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part. The device especially utilises the non-pulsatile element of a patient's blood. There is also described a method of measuring blood glucose levels and a device programmed so as to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

NON-INVASIVE MEASUREMENT OF BLOOD ANALYTES

This invention relates to a novel monitor, particularly a monitor for the non-invasive measurement of glucose in eg diabetics and a method for determining glucose levels.

Diabetes mellitus (abbreviated to diabetes) is the name for a group of chronic or lifelong diseases that affect the way the body uses food to make energy necessary for life. Primarily, diabetes is a disruption of carbohydrate (sugar and starch) metabolism and also affects fats and proteins. In people who have diabetes the glucose levels vary considerably being as high as 40 mmol/l and as low as 2 mmol/l. Blood glucose levels in people without diabetes vary very little, staying between 3 and 7 mmol/l. These levels follow the typical patterns shown in Figure 1a.

Hyperglycaemia (high blood glucose)

Both insulin dependant diabetes (IDDM) and non-insulin dependant diabetes (NIDDM) are associated with serious tissue complications which characteristically develop after 10-20 years duration of diabetes. Diabetic eye disease, retinopathy, is the commonest cause of blindness in western countries in people under the age of 65 years. Diabetic renal disease, nephropathy, is an important cause of kidney failure in the community. Diabetic neuropathy affects the peripheral nerves causing impaired sensation and leg ulcers, and damage to the autonomic nervous system causes postural hypertension (low blood pressure on standing) and diarrhoea. Atherosclerosis is 2-4 times as high in diabetic as non-diabetic people and manifest as an increased frequency of myocardial infarction (heart attacks), cerebrovascular disease (strokes) and the peripheral vascular disease (causing reduced circulation to the limbs and the risk of gangrene and amputation).

For many years it has been something of an article of faith in clinical diabetes that the cause of the complications is exposure of the tissues over many years to the higher than normal blood glucose levels which have been usual in most treated diabetic patients. Conclusive proof of this theory has only recently become

available; the landmark Diabetes Control and Complications Trial (DCCT) in North America was announced in 1993 and showed that IDDM patients randomly assigned to an intensive and optimised insulin treatment programme designed to produce near-normal blood glucose levels had significantly less retinopathy, kidney disease and neuropathy over a 9-year period that patients assigned to ordinary treatment (ie poor control).

The DCCT has been a major stimulus to physicians around the world to renew efforts to improve control in diabetic patients, and to develop improved methods of obtaining good control and of monitoring these patients.

Hypoglycaemia (low blood glucose)

An important additional finding in the DCCT was that the frequency of hypoglycaemia was three-fold higher in the well-controlled patients than those with ordinary control. This confirms the long-standing appreciation by physicians that hypoglycaemia is extremely frequent in IDDM, and especially so in those that are well controlled. There are many reasons for this including mistiming of insulin injections and food, erratic absorption of insulin, and impaired secretion in some diabetic patients of the so-called counter regulatory hormones such as adrenaline and glucagon that oppose the action of insulin.

About one third of IDDM patients have no warning symptoms of hypoglycaemia (eg sweating, nausea, blurred vision, palpitations) and they rely on intermittent self-monitoring of blood glucose to detect dangerously low glucose levels. The consequences of hypoglycaemia include impaired cognition and consciousness, and eventually coma.

Since the late 1970's, an increasing number of diabetic patients, mostly IDDM, have been measuring their own blood glucose concentrations using finger-prick capillary blood samples. Self blood glucose monitoring (SBGM) is used by diabetics in the home to detect hypoglycaemia or hyperglycaemia and take corrective action such as

taking extra food to raise the blood glucose or extra insulin to lower the blood glucose. The measurements, which are made using a low-cost, hand-held blood glucose monitor (BGM), also allow the physician to adjust the insulin dosage at appropriate times so as to maintain near normoglycaemia.

BGMs use either reflectance photometry or an electrochemical method to measure the glucose concentration. Reflectance photometry measures the amount of light reflected from the reagent-impregnated test strip that has reacted with a drop of blood. The operator pricks the finger of the patient or earlobe with a sterile lancet or uses anticoagulated whole blood collected in heparin and then places the blood on the test strip. The operator must place the blood onto the test strip at the time the monitor begins its timing sequence. This step is critical because under-timing (under-incubation) or over-timing (over-incubation) of the reaction may cause inaccurate measurements. At the audible signal, the operator wipes or blots the excess blood off the outside of the test strip. The operator then inserts the strip into the monitor for measurement.

In the electrochemical method a disposable single-use enzyme electrode test strip is used. When the test specimen is placed onto the test strip, an enzymatic reaction occurs that results in a current through the strip. The current is directly proportional to the concentration of glucose in the specimen.

The main disadvantages of SBGM systems are poor patient acceptance because the technique is painful, only intermittent assessment of diabetic control is possible and readings during the night or when the patient is otherwise occupied such as during driving are not possible. It is estimated that less than half of the IDDM patients in the US perform SBGM.

Further, glucose values obtained with BGMs may not agree with clinical laboratory results. Routine laboratory measurements of glucose are performed on either serum

or plasma venous blood specimens that correspond with glucose concentrations measured on whole blood glucose analysers.

Whole blood glucose values are lower than those obtained from either serum or plasma. Although glucose is not a static component in human blood, changes in blood glucose concentration following food intake in normal and hyperglycaemic conditions are reasonably predictable. Similarly, the variation in glucose concentration as blood passes from arteries or capillaries to veins has also been documented. Therefore, over time, repeated measurement of blood glucose from the same patient may diverge widely. Also, blood obtained simultaneously by finger stick and venipuncture may not have the same glucose concentration. (Venous blood may contain 1 mmol/l less glucose than capillary blood if the same samples are obtained within 1-2 hours after carbohydrate intake).

Furthermore, the haematocrit of the patient (the volume of cells, mostly erythrocytes, expressed as a percentage of the volume of whole blood in a sample) influences glucose values, and whole blood glucose measurements must be corrected for this. Unfortunately, because BGMs cannot automatically correct for the haematocrit of the patient, an error of 5-15% may be introduced.

There is widespread agreement that for self-monitoring in the home the reluctant acceptance of the current finger-stick method is the main reason why the development of a non-invasive measurement technique has such high priority.

A non-invasive measurement device is known from US Patent No 5,553,613. US '613 describes a technique which uses the pulsatile component of the light intensity transmitted through the finger, from which to derive the glucose concentration non-invasively. It does this by using the wavelengths 805nm, 925nm, 970nm and the range 1000-1100nm. The measurements were made by transmission, ie light was passed through the finger. However, as mentioned above, US '613 specifically relies upon the pulsatile component of the light transmitted through the patient. Such a

pulsatile technique has clear disadvantages in that the pulsatile component of the light signal, whether transmitted or reflected, is less than 2% of the total signal. Thus, prior art devices which use only the pulsatile component are much less sensitive and much more vulnerable to patient movement which can cause interference which is in the order of a few hundred times the relevant signal.

Moreover, the pulsatile signal identifies arterial blood specifically. Whilst this is advantageous when considering the pulmonary circulation of a patient, it provides no information on the patient's systemic circulation which is important for glucose determination. Further, pulsatile techniques are limited to use on body extremities, eg finger, ear lobe or the ball of the foot in babies or neo-nates.

The present invention overcomes or mitigates the disadvantages of the prior art by using a plurality of closely associated transmitters and generators which allows an "average" evened out signal to be produced and is capable of utilising the non-pulsatile element of the patient's blood flow. In addition, a further advantage of the invention is that it allows the measurement of oxygen saturation (SO_2). Furthermore, the invention permits the measurement of haemoglobin index (HbI) and/or temperature.

According to the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part.

A preferred embodiment of the invention is one wherein the non-pulsatile element of the patient's blood is utilised. Thus, according to a further feature of the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part as herein before described which is adapted to measure the non-pulsatile element of a patient's blood. In a further preferred embodiment the

device measures the pulsatile and non-pulsatile elements of a patient's blood. The device may be so adapted by being provided with a plurality of closely associated transmitters and generators which allows an "average" evened out signal to be produced.

Although various analytes may be measured, the detector of the invention is especially useful in measuring blood glucose level. We especially provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part as herein before described which is adapted to measure blood glucose levels.

The device may be capable of measuring other parameters either separately or in addition to blood glucose. An especially advantageous feature is the device may be adapted to measure blood oxygen saturation (SO_2).

As a further preferred embodiment we provide a device which is adapted to measure the haemoglobin index (HbI) and/or temperature of a patient's blood.

The device may be adapted for use, with any body part although it is preferable that it can be a finger or thumb.

The number of transmitter fibres may vary although we have found that 18 transmitter fibres works well. The number of detector fibres may be the same or different to the number of transmitter fibres, but may vary and we have found that 12 detector fibres works well. The diameter of the detector and the transmitter fibres may be the same or different and may vary, a diameter of 250μm is preferred.

The detector fibres are preferably positioned to detect reflected light rather than transmitted light.

The wavelength used in the transmitter fibres will generally be from 500 to 1100nm. However, it is a further feature of the invention to provide a detector as hereinbefore

described which also measures haemoglobin index (HbI) and/or oxygen saturation (SO_2) of blood. For such measurement, specific wavelengths are used, namely 500.9nm, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm. The preferred wavelengths for measuring blood glucose are from 800nm to 1100nm.

According to a further feature of the invention we provide a method of measuring blood glucose levels which comprises placing a non-invasive measuring device as hereinbefore described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part.

In a yet further feature of the invention we provide a device according to as herein before described programmed so as to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation. Clearly, since blood oxygen saturation is dependent upon both the haemoglobin index and the oxygen index, the computer is programmed so as to calculate these equations first if blood oxygen saturation is to be calculated.

We also provide a computer programme product comprising a computer readable medium having thereon computer programme code means, when said programme is loaded, to make the computer execute a procedure to calculate one or more of the haemoglobin index, the oxygen index and the "whole blood" oxygen saturation as herein before described.

The invention will now be illustrated but in no way limited by reference to the following example and drawings in which;

Figure 1 is a plot of the predicted glucose values against the measured glucose values; and

Figure 1a is a graph comparing normal blood glucose levels with those of a diabetic.

Example 1

Glucose measurement

In vivo measurements using the MCPD spectrophotometer were carried out at 10 min intervals on the fingertips of 8 volunteers during the course of glucose tolerance tests and the results compared with those measured using a conventional blood glucose monitor. In addition, parallel measurements of local blood flow (laser Doppler flux) and temperature were made.

The analysis which is presented here uses the same wavelength range used in the previous glucose studies carried out namely: 805nm, 925nm, 970nm and the broadband average 1000-1100nm, but additionally wavelengths sampled at regular intervals in the entire range 800nm to 1100nm. Intervals of 1.96nm worked well.

Earlier work demonstrated that the glucose-dependent signal emanates from haemoglobin. Furthermore, although the 805nm wavelength could be used to compensate for small changes in haemoglobin concentration large changes continued to interfere with the sensitivity for glucose. It was furthermore recognised that changes in haemoglobin oxygenation would cause absorption changes from 800nm to 1100nm. As in all physiological measurements carried out in the peripheral circulation, temperature is also likely to be a controlling parameter. In the novel analysis carried out on the intensity spectra in the experiments carried out here, the three parameters haemoglobin concentration, oxygen saturation and temperature were introduced into the multiple linear regression analysis along with the near infrared parameters previously used.

Experimental

13 glucose tolerance tests (GTTs) were carried out on 8 different volunteers. In one case, 200ml water was given instead of the solution of 75g glucose in 200ml water, a real GTT was subsequently carried out on the same volunteer. In one volunteer five GTTs were carried out on separate occasions. One volunteer had diabetes.

All measurements were carried out with an Otsuka Optronics Photal MCPD-1000 photodiode array lightguide spectrophotometer. The 0.2mm slit was used for the diffraction grating giving a full width at half maximum transmission of 7.2nm, comparable with the glucose monitor. Using the supplied software, the instrument allows access to data points at 1.94nm intervals within the wavelength range 300-1100nm. The range displayed during the glucose experiments was 500-1100nm. In order to mimic the broad bandwidth characteristics of the previous glucose monitor above 1000nm, all measurements were averaged over the range 1000-1100nm. Quartz lightguides were used in conjunction with a 400W quartz-halogen light source.

A lightguide bundle, which consisted of 18 transmitting and 12 receiving fibres each of 250μm diameter, was attached to the fingertip of the subject by means of a laser Doppler probe holder. Recordings of spectra were made at 10 min intervals throughout the test using the MCPD spectrophotometer described above. These recordings were accompanied by parallel measurements of glucose concentrations in blood, obtained by pinprick of a contralateral finger with the aid of a Softelix pro lancet system, using a Boehringer Manheim Advantage® glucose monitor. The lightguide was removed from the finger after each measurement and new dark and reference spectra recorded before each new measurement. A total of 13 measurements were carried out over a 2 hour period.

Careful selection of integrating time and the intensity of the reference spectrum enabled the simultaneous record of spectra that covered not only the range 800-1100nm, but also the visible range from 500-600nm. This enabled the evaluation of skin haemoglobin saturation (SO_2) and haemoglobin concentration (HbI) (Harrison DK *et al.* (1993) *Phys Meas* **14**: 241-52) from the same spectra as those being analysed for glucose (see below).

A Moor Instruments DRT4 laser Doppler perfusion monitor was used to measure blood flow changes in the adjacent finger. The probe incorporated a thermal sensor, which was used to measure skin temperature (note: also on the adjacent finger) throughout the experiment.

Derivation of HbI and SO_2

HbI and SO_2 were derived from the absorption spectra measured from 500.8 to 586.3nm using a computer program VOXYG written for the purpose. The program carried out the following calculations.

Haemoglobin Index

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100$$

Oxygenation Index

$$OXI=(c-d)/11.7-(d-c)/11.6)*100/HbI$$

Oxygen Saturation

$$SO_2=100*(OXI+0.43)/1.5$$

where a = absorption value at 500.9nm
b = absorption value at 528.1nm
c = absorption value at 549.5nm
d = absorption value at 561.1nm
e = absorption value at 572.7nm
f = absorption value at 586.3nm

MULTIPLE LINEAR REGRESSION ANALYSIS

A data file A was created containing the full absorption spectral data (800-1100nm in 1.96nm steps) from all 12 GTTs in the series. The absorption values in the file are defined in "absorption units" referred to here as ABUs. The other data contained in the file were time, experiment identification, glucose concentration (invasive), Hbl, SO_2, temperature, and laser Doppler flux.

A number of secondary files were created whereby a sequence of "normalisations" of the data were performed:

- B - ABU data of A was normalised by subtraction of the absorption of the values at 802nm (ie $ABU_A - ABU_{802}$). This is similar to the way in which previous data was treated.
- C - ABU data of B was further normalised by division by the Hbl value (ie ABU_B/Hbl). This was designed to take into account of the results of the *in vitro* experiments which showed that normalisation at, then, 805nm did not fully compensate for changes in haemoglobin concentration.
- D - ABU data of C was further normalised by division by the SO_2 value (ie ABU_C/SO_2) to take into account the influence of changes in the relative concentrations of oxygenated haemoglobin (HbO_2) and deoxygenated haemoglobin (Hb) on the infrared spectrum.
- E - SBU data of D was yet further normalised by subtraction of the value at the assumed water peak (ie $ABU_D - ABU_{949}$) in an attempt to take into account changes in water content.

The types and orders of normalisations may vary, and the above are examples.

The above files were then subjected to multiple linear regression, analysis using SPSS for Windows 6.1.2. All of the wavelengths available in the above data files, ie 800nm to 1100nm in 1.96nm steps were entered as independent variables. The results of the multiple wavelength regressions are given below. The regressions

include only the spectral data and not HbI, SO_2 or temperature as further independent variables at this stage.

	r	Standard Error (SE) (mM)	No of Wavelengths Included
A	0.48	2.81	4
B	0.89	1.69	37
C	0.80	2.05	21
D	0.89	1.61	31
E	0.93	1.40	48

The predicted values from the last correlation using data file E are plotted against the measured glucose values in Figure 1. The predicted values are given as standardised to the mean and number of standard deviations on the left hand side of the y-axis and as mM on the right hand side.

The results obtained using the multi-wavelength analysis are significant improvements to those using the original parameters applied to the collective results. Figure 1 could indicate that the method may eventually allow a universal calibration, or at least one based on a particular individual, particularly if the ways in which the spectra are normalised are varied.

Above multiple regression analyses result in regression equations whose coefficients can be incorporated into an equation to produce a new parameter "calculated Glucose". This, together with the parameters HbI, SO_2 and temperature can then be incorporated into a further regression equation for each individual GTT.

Least squares fitting of mean "calibration spectra" recorded from the GTT series could be used for a universal or individual calibration.

CLAIMS

1. A device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part.

2. A device according to Claim 1 characterised in that it is adapted to utilise the non-pulsatile element of a patient's blood.

3. A device according to Claim 1 characterised in that it is adapted to utilise the non-pulsatile and the pulsatile elements of a patient's blood.

4. A device according to Claim 2 characterised in that it is so adapted by being provided with a plurality of closely associated transmitters and generators which allow an "average" evened out signal to be produced.

5. A device according to Claim 1 characterised in that it is adapted to measure blood glucose levels.

6. A device according to Claim 1 characterised in that it is adapted to measure blood oxygen saturation (SO_2).

7. A device according to Claim 1 characterised in that it is adapted to measure the haemoglobin index (HbI).

8. A device according to Claim 1 characterised in that it is adapted to measure the temperature of a patient's blood.

9. A device according to Claim 1 characterised in that it is adapted to measure two or more analytes in blood, which analytes are selected from blood glucose levels, blood oxygen saturation (SO_2), the haemoglobin index (HbI) and the temperature of a patient's blood.

10. A device according to Claim 9 characterised in that it is adapted to measure blood glucose levels, blood oxygen saturation (SO_2), the haemoglobin index (HbI) and the temperature of a patient's blood.

11. A device according to Claim 1 characterised in that it is adapted to measure of one or more analytes in blood in a patient's finger or thumb.

12. A device according to Claim 1 characterised in that it is provided with a greater number of transmitter fibres than detector fibres.

13. A device according to Claim 1 characterised in that it is provided with from 12 to 24 transmitter fibres.

14. A device according to Claim 13 characterised in that it is provided with 18 transmitter fibres.

15. A device according to Claim 1 characterised in that it is provided with from 6 to 18 detector fibres.

16. A device according to Claim 14 characterised in that it is provided with 12 detector fibres.

17. A device according to Claim 1 characterised in that diameter of the fibres is from 200 - 300μm.

18. A device according to Claim 1 characterised in that diameter of the fibres is 250µm.

19. A device according to Claim 1 characterised in that the detector fibres are positioned to detect transmitted light rather than reflected light.

20. A device according to Claim 1 characterised in that the wavelength used in the transmitter fibres is from 500 to 1100nm.

21. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is from 800 to 1100nm.

22. A device according to Claim 21 characterised in that the wavelength used in the transmitter fibres is 805nm, 925nm, 970nm and the broadband average 1000-1100nm.

23. A device according to Claim 21 characterised in that the wavelengths are sampled at regular intervals of 1.96nm in the entire range 800nm to 1100nm.

24. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is from 500 to 600nm.

25. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is 500.9nm, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm.

26. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is 500.9nm, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm; and from 800nm to 1100nm.

27. A method of measuring blood glucose levels which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part.

28. A method according to Claim 27 characterised in that the non-pulsatile element is used.

29. A method according to Claim 28 characterised in that the non-pulsatile and pulsatile element is used.

30. A device according to Claim 1 programmed so as to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

31. A device according to Claim 30 programmed so as to conduct a multiple linear regression analysis on spectral data collected by the detectors.

32. A device according to Claim 30 wherein the Haemoglobin Index is calculated using the equation:

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100$$

where a = absorption value at 500.9nm
b = absorption value at 528.1nm
c = absorption value at 549.5nm
e = absorption value at 572.7nm.

33. A device according to Claim 30 wherein the Oxygenation Index is calculated using the equation:

$$OXI=(ee-d)/11.7-(d-c)/11.6)*100/HbI$$

where c = absorption value at 549.5nm

d = absorption value at 561.1nm

e = absorption value at 572.7nm.

34. A device according to Claim 30 wherein the Oxygen Saturation (SO_2) is calculated using the equation:

$$SO_2=100*(OXI+0.43)/1.5.$$

35. A method of measuring blood oxygen saturation which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part using a device according to Claim 34.

36. A computer programme product comprising a computer readable medium having thereon computer programme code means, when said programme is loaded, to make the computer execute a procedure to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

37. A computer programme according to Claim 36 wherein the computer programme code means will make the computer execute a procedure to calculate one or more of :

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100;$$

$$OXI=(e-d)/11.7-(d-c)/11.6)*100/HbI; \text{ and}$$

$$SO_2=100*(OXI+0.43)/1.5$$

where a = absorption value at 500.9nm

b = absorption value at 528.1nm

c = absorption value at 549.5nm

d = absorption value at 561.1nm

e = absorption value at 572.7nm

38. A device substantially as described with reference to the accompanying examples and drawings.

P36002WO.1

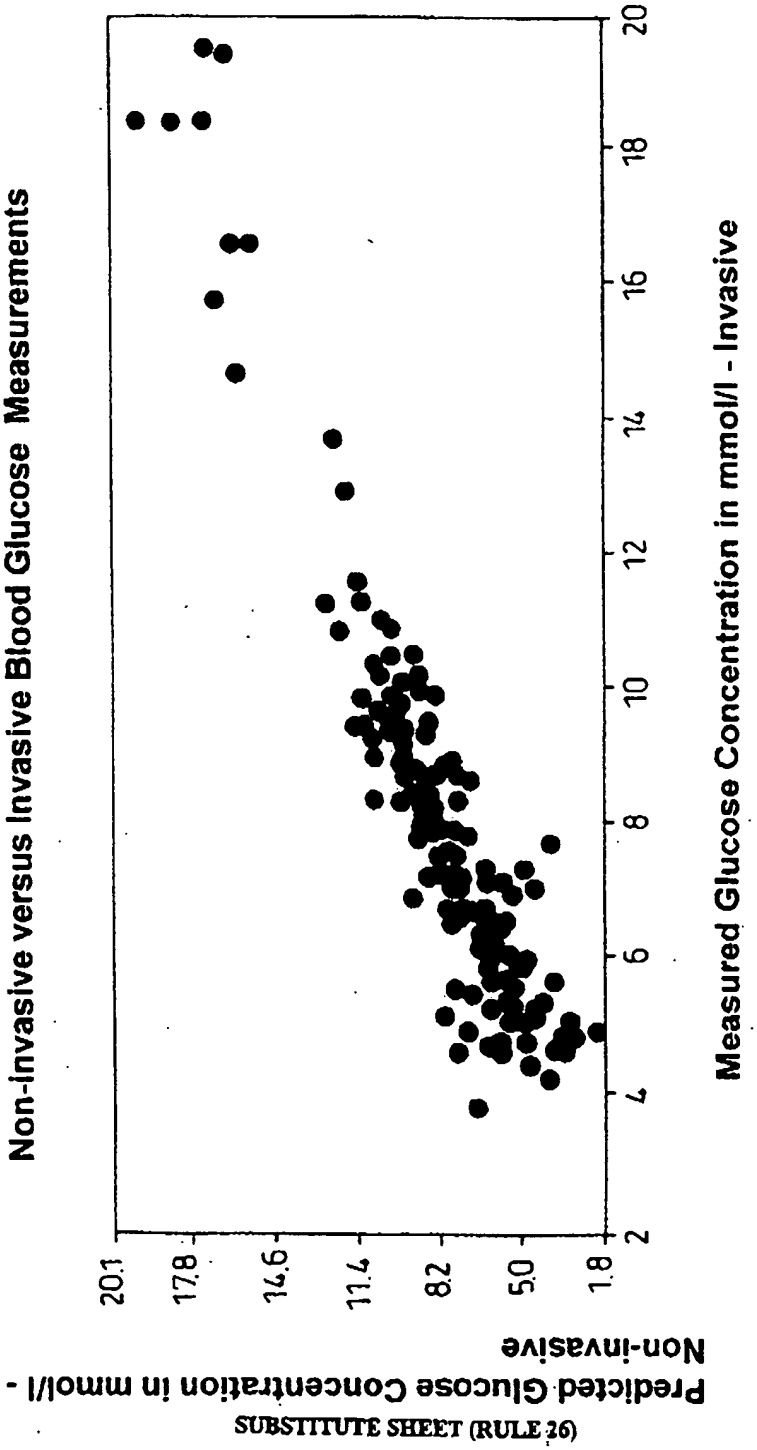
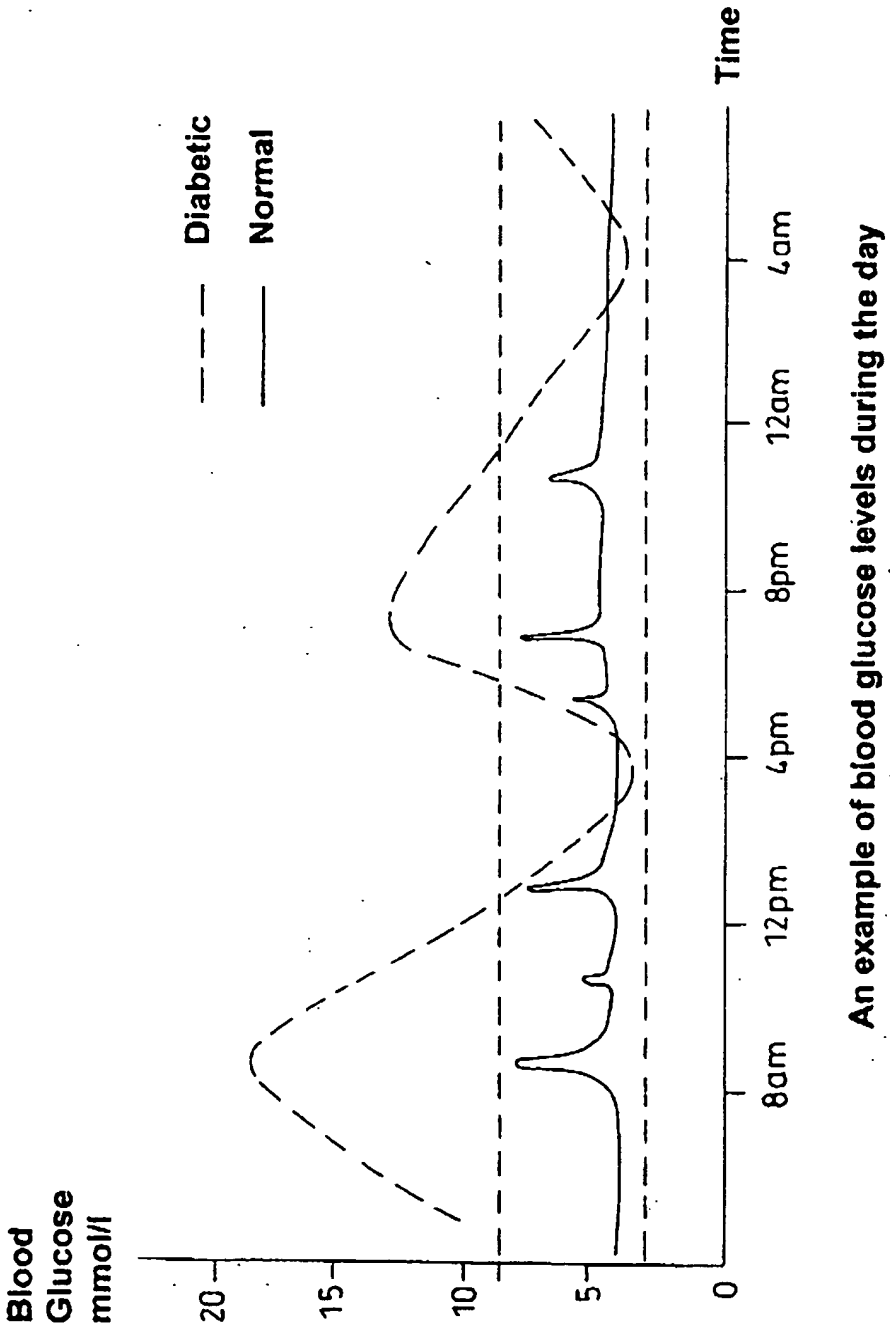

Non-invasive versus Invasive Blood Glucose Measurements
Predicted Glucose Concentration in mmol/l - Non-invasive
Measured Glucose Concentration in mmol/l - Invasive
20.1
17.8
14.6
11.4
8.2
5.0
1.8
2
4
6
8
10
12
14
16
18
20

Fig. 1



An example of blood glucose levels during the day

Fig. 1a

INTERNATIONAL SEARCH REPORT

Interr nal Application No
PCT/GB 99/02127

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61B5/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 755 226 A (3M) 26 May 1998 (1998-05-26) the whole document	1-4,6,7, 9-11,15, 19-22, 24-31,38
X	WO 97 27800 A (DIASENSE) 7 August 1997 (1997-08-07) the whole document	1,5,12, 15-17

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
27 September 1999	05/10/1999

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Lemercier, D

Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

I National application No.
PCT/GB 99/02127

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. [X] Claims Nos.: 36,37
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Program for computers

2. [ ] Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. [ ] Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. [ ] As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. [ ] As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. [ ] No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

..formation on patent family members

Inter ...nal Application No
PCT/GB 99/02127

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5755226 A	26-05-1998	US 5553615 A	10-09-1996
		EP 0742896 A	20-11-1996
		JP 9508291 T	26-08-1997
		WO 9520757 A	03-08-1995
WO 9727800 A	07-08-1997	AU 1846897 A	22-08-1997
		EP 0889703 A	13-01-1999

Form PCT/ISA/210 (patent family annex) (July 1992)

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
SPG/P36002WO		
International application No.	International filing date (day/month/year)	Priority date (day/month/year)
PCT/GB99/02127	02/07/1999	04/07/1998

International Patent Classification (IPC) or national classification and IPC
A61B5/00

Applicant
WHITLAND RESEARCH LIMITED et al.

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 7 sheets, including this cover sheet.

   ☒ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of 7 sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☐ Certain documents cited
   - VII ☐ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
01/02/2000	13.10.2000
Name and mailing address of the international preliminary examining authority: European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Fontenay, P Telephone No. +49 89 2399 2646

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/02127

I. Basis of the report

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

Description, pages:

1-4,6-12	as originally filed	
5,5a	with telefax of	01/08/2000

Claims, No.:

1-32	with telefax of	01/08/2000

Drawings, sheets:

1/2,2/2	as originally filed

2. The amendments have resulted in the cancellation of:

- ☐ the description, pages:
- ☐ the claims, Nos.:
- ☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

- ☐ the entire international application.
- ☒ claims Nos. 7,32.

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/02127

because:

☐ the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (*specify*):

☒ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. 7 32 are so unclear that no meaningful opinion could be formed (*specify*):

see separate sheet

☐ the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.

☐ no international search report has been established for the said claims Nos. .

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims 2-6, 8-22, 27-31
	No:	Claims 1, 23-26
Inventive step (IS)	Yes:	Claims 27-31
	No:	Claims 2-6, 8-22
Industrial applicability (IA)	Yes:	Claims 1-32
	No:	Claims

2. Citations and explanations

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

INTERNATIONAL PRELIMINARY EXAMINATION REPORT - SEPARATE SHEET International application No. PCT/GB99/02127

Re Item III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The subject-matter of claims 7 and 32 is not clearly defined (Article 6 PCT).

III.1 The wording "characterized in that it is adapted to measure the temperature of a patient's blood" as it appears in claim 7 is ambiguous. It could suggest that the temperature is obtained from the measurements carried out within the claimed device in the same way as it calculates the haemoglobin index or the blood oxygen saturation or that it contains a thermal sensor as it is suggested on page 10, lines 2-4 of the description.

III.2 The claims should not refer to the drawings or to parts of the description (Rule 6.2 (a) PCT). Claim 32 is accordingly not allowable.

Re Item V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1 = WO-A-9727800
D2 = US-A-5755226

V.1 The subject-matter of independent claim 1 is not new in the sense of Article 33(2) PCT.

D2 discloses a device for the non-invasive measurement of one or more analyte in blood (haematocrit) in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibers positioned to transmit light to the body part (see D2, column 15, line 44 - column 16, line 17, figure 4) and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part (see D2, column 13, lines 47-62). Moreover, the device disclosed in D2 is adapted to utilise the non-pulsatile element of a patient's blood (see D2, column 6, lines 27-47; column 8, lines 53 -

column 9, line 27). It is here noted that the fact that the claimed device is adapted to utilise the non-pulsatile element of a patient's blood does not exclude that it utilises additional components.

It follows that all the features of claim 1 are known in combination from D2. The subject-matter of claim 1 is accordingly not new.

The device of D2 is programmed so as to calculate a multiplicity of analytes concentration and in particular haematocrit and oxygen saturation. Claim 25 is accordingly also not new. It is also proposed in D1 to base the analysis on multiple linear regression so that claim 26 is also not new.

V.2 The features of dependent claims 2-6, 8-18 and 20-22 are known from or rendered obvious by the prior art as illustrated by documents D1 or D2 and as indicated in the International Search Report in relation with original claims 3-7, 9-12, 15, 17, 19, 21, 22 and 24-26. The dependent claims 2-6, 8-17, and 20-22 therefore do not appear,to contain any additional features which, in combination with the features of claim 1 to which they refer, would involve an inventive step.

It is in particular noted that in D2, a plurality of associated transmitters and generators are used and that an average signal is produced so as to consider the non-pulsatile component of the signal (see D2, column 23, line 1- column 24, line 53).

V.3 The feature of the sampling interval of 1,96 nm as it appears in claim 19, constitutes a constructional detail as to the device in order to obtain the information over the whole spectra. This feature cannot justify a positive inventive step assessment since the skilled man will arrive at something falling under the terms of the claims by the exercise of routine trials according to the intended purpose i.e. in order to obtain sufficient information over said spectra. The applicant may also refer to the Guidelines, PCT/GL/3 Chapter IV, 8.8 (C1)(ii).

V.4 The subject-matter of claims 23 and 24, as may be understood (see comments under point VIII) is already known from the prior art. Document D2 discloses a method of measuring blood glucose which comprises placing a non-invasive

measuring device as claimed against a body part (a finger) of a patient and using the detector to measure the light reflected from the body part (see D2, figures 2. 4, 7). In D2, as explained above under section V.1, the non-pulsatile component of the signal is utilised.

The subject-matter of claims 23 and 24 is therefore not new considering the teaching of D2.

V.5 The subject-matter of claim 27 and 31 is not disclosed as such in D2. None of the documents cited in the search report suggest to calculate the Haemoglobin index on the basis of the equation indicated in claim 27 or 31. The subject-matter of claim 27 is accordingly considered to be new and inventive in the sense of article 33(2) and 33(3) PCT considering the prior art as presently known.

Claim 28, when depending on claim 27 (see comments under point VIII.3) would accordingly also be new and inventive. The same would apply to claim 29 when depending on claim 28 (see the comments under point VIII.3).

V.6 The method defined in claim 30 seems to refer to the use of a device as defined in claim 29 (see comments under section VIII). The device of claim 29 being considered as new and inventive, when depending on claim 28, the same applies to said use.

Re Item VIII Certain observations on the international application

VIII.1 The subject-matter of claim 1 is not clearly defined (Article 6 PCT). The wording in the characterising part of claim 1: "is adapted to utilise the non-pulsatile element of a patient's blood is too vague because of the terms "adapted to utilise" which presents a broad meaning. It is in particular considered that any device which senses a signal with a pulsatile and a non-pulsatile component may be considered as being adapted to utilise the non-pulsatile component. The present meaning does not permit to "define" the matter for which protection is sought as requested under Article 6.

VIII.2 The subject-matter of claims 23 and 30 is not clearly defined (Article 6 PCT) because it refers to the device as "herein before described". Said wording is not allowable in the claims. It has been assumed in section V above, that the wording "measuring device according to any of the preceding claims" had been intended.

Claim 30 is also not clear because it refers to a first device (as herein before described) and then to a device according to claim 29.

VIII.3 Claim 28 is not clear because it refers to claim 30. Moreover, claim 28 refers to the parameter HBI which is defined in claim 27. The same applies to claim 29 which refers to the parameter SO_2 which is defined in claim 28. It has accordingly been assumed in the present report that claim 28 refers to claim 27 and that claim 29 refers to claim 28.

pulsatile technique has clear disadvantages in that the pulsatile component of the light signal, whether transmitted or reflected, is less than 2% of the total signal. Thus, prior art devices which use only the pulsatile component are much less sensitive and much more vulnerable to patient movement which can cause interference which is in the order of a few hundred times the relevant signal.

Moreover, the pulsatile signal identifies arterial blood specifically. Whilst this is advantageous when considering the pulmonary circulation of a patient, it provides no information on the patient's systemic circulation which is important for glucose determination. Further, pulsatile techniques are limited to use on body extremities, eg finger, ear lobe or the ball of the foot in babies or neo-nates.

International Patent Application No. WO 97/27800 discloses a device for the non-invasive measurement of blood analytes using light transmitted through or reflected from a body part.

However, the invention disclosed in the prior art alsosuffers from the disadvantage that, *inter alia*, only the pulsatile element of the transmitted/reflected signal is exploited.

US Patent No. 5,755,226 describes an apparatus for the non-invasive measurement of blood glucose levels. Furthermore, the disclosed invention does not describe the utilisation of the non-pulsatile element of a patient's blood in the determination of blood analytes.

The present invention overcomes or mitigates the disadvantages of the prior art by using a plurality of closely associated transmitters and generators which allows an "average" evened out signal to be produced and is capable of utilising the non-pulsatile element of the patient's blood flow. In addition, a further advantage of the invention is that it allows the measurement of oxygen saturation (SO_2).

Furthermore,the invention permits the measurement of haemoglobin index (HbI) and/or temperature.

According to the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part.

A preferred embodiment of the invention is one wherein the non-pulsatile element of the patient's blood is utilised. Thus, according to a further feature of the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part as herein before described which is adapted to measure the non-pulsatile element of a patient's blood. In a further preferred embodiment the

CLAIMS

1. A device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part characterised in that it is adapted to utilise the non-pulsatile element of a patient's blood.

2. A device according to Claim 1 characterised in that it is adapted to utilise the non-pulsatile and the pulsatile elements of a patient's blood.

3. A device according to Claim 2 characterised in that it is so adapted by being provided with a plurality of closely associated transmitters and generators which allow an "average" evened out signal to be produced.

4. A device according to Claim 1 characterised in that it is adapted to measure blood glucose levels.

5. A device according to Claim 1 characterised in that it is adapted to measure blood oxygen saturation (SO_2).

6. A device according to Claim 1 characterised in that it is adapted to measure the haemoglobin index (Hbl).

7. A device according to Claim 1 characterised in that it is adapted to measure the temperature of a patient's blood.

8. A device according to Claim 1 characterised in that it is adapted to measure two or more analytes in blood, which analytes are selected from blood glucose levels,

AMENDED SHEET

blood oxygen saturation (SO_2), the haemoglobin index (Hbl) and the temperature of a patient's blood.

9. A device according to Claim 8 characterised in that it is adapted to measure blood glucose levels, blood oxygen saturation (SO_2), the haemoglobin index (Hbl) and the temperature of a patient's blood

10. A device according to Claim 1 characterised in that it is adapted to measure of one or more analytes in blood in a patient's finger or thumb.

11. A device according to Claim 1 characterised in that it is provided with a greater number of transmitter fibres than detector fibres.

12. A device according to Claim 1 characterised in that it is provided with from 6 to 18 detector fibres.

13. A device according to Claim 12 characterised in that it is provided with 12 detector fibres.

14. A device according to Claim 1 characterised in that diameter of the fibres is from 200 - 300μm.

15. A device according to Claim 1 characterised in that the detector fibres are positioned to detect transmitted light rather than reflected light.

16. A device according to Claim 1 characterised in that the wavelength used in the transmitter fibres is from 500 to 1100nm.

17. A device according to Claim 16 characterised in that the wavelength used in the transmitter fibres is from 800 to 1100nm.

AMENDED SHEET

18. A device according to Claim 17 characterised in that the wavelength used in the transmitter fibres is 805nm, 925nm, 970nm and the broadband average 1000-1100nm.

19. A device according to Claim 17 characterised in that the wavelengths are sampled at regular intervals of 1.96nm in the entire range 800nm to 1100nm.

20. A device according to Claim 16 characterised in that the wavelength used in the transmitter fibres is from 500 to 600nm.

21. A device according to Claim 16 characterised in that the wavelength used in the transmitter fibres is 500.9nm, 528.1nm, 549.5nm, 561.1um, 572.7nm and 586.3nm.

22. A device according to Claim 16 characterised in that the wavelength used in the transmitter fibres is 500.9um, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm; and from 800nm to 1100nm.

23. A method of measuring blood glucose levels which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part characterised in that the non-pulsatile element is used.

24. A method according to Claim 23 characterised in that the non-pulsatile and pulsatile element is used.

25. A device according to Claim 1 programmed so as to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

26. A device according to Claim 25 programmed so as to conduct a multiple linear regression analysis on spectral data collected by the detectors.

AMENDED SHEET

27. A device according to Claim 25 wherein the Haemoglobin Index is calculated using the equation:

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100$$

where a = absorption value at 500.9nm
b = absorption value at 528.1nm
c = absorption value at 549.5nm
e = absorption value at 572.7nm
f = absorption value at 586.3nm.

28. A device according to Claim 30 wherein the Oxygenation Index is calculated using the equation:

$$OXI=(ee-d)/11.7-(d-c)/11.6)*100/HbI$$

where c = absorption value at 549.5nm
d = absorption value at 561.1nm
e = absorption value at 572.7nm.

29. A device according to Claim 25 wherein the Oxygen Saturation (SO_2) is calculated using the equation:

$$SO_2=100*(OXI+0.43)/1.5.$$

30. A method of measuring blood oxygen saturation which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part using a device according to Claim 29.

AMENDED SHEET

31. A computer programme product comprising a computer readable medium having thereon computer programme code means, when said programme is loaded, to make the computer execute a procedure to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation characterised in that the computer programme code means will make the computer execute a procedure to calculate one or more of :

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100;$$

$$OXI=(e-d)/11.7-(d-c)/11.6)*100/HbI; \text{ and}$$

$$SO_2=100*(OXI+0.43)/1.5$$

where a = absorption value at 500.9nm
b = absorption value at 528.1nm
c = absorption value at 549.5nm
d = absorption value at 561.1nm
e = absorption value at 572.7nm
f = absorption value at 586.3nm

32. A device substantially as described with reference to the accompanying examples and drawings.

PJ6002WO.3claims

AMENDED SHEET

INTERNATIONAL SEARCH REPORT

Interr ɔnal Application No
PCT/GB 99/02127

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61B5/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 755 226 A (3M) 26 May 1998 (1998-05-26) the whole document ---	1-4,6,7, 9-11,15, 19-22, 24-31,38
X	WO 97 27800 A (DIASENSE) 7 August 1997 (1997-08-07) the whole document -----	1,5,12, 15-17

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

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"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
27 September 1999	05/10/1999

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl, Fax: (+31–70) 340–3016	Lemercier, D

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Form PCT/ISA/210 (second sheet) (July 1992)

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB 99/02127

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

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because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Program for computers

2. [ ] Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

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Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

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2. [ ] As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

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Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

..formation on patent family members

Inter...nal Application No
PCT/GB 99/02127

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5755226 A	26-05-1998	US 5553615 A	10-09-1996
		EP 0742896 A	20-11-1996
		JP 9508291 T	26-08-1997
		WO 9520757 A	03-08-1995
WO 9727800 A	07-08-1997	AU 1846897 A	22-08-1997
		EP 0889703 A	13-01-1999

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CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

pulsatile technique has clear disadvantages in that the pulsatile component of the light signal, whether transmitted or reflected, is less than 2% of the total signal. Thus, prior art devices which use only the pulsatile component are much less sensitive and much more vulnerable to patient movement which can cause interference which is in the order of a few hundred times the relevant signal.

Moreover, the pulsatile signal identifies arterial blood specifically. Whilst this is advantageous when considering the pulmonary circulation of a patient, it provides no information on the patient's systemic circulation which is important for glucose determination. Further, pulsatile techniques are limited to use on body extremities, eg finger, ear lobe or the ball of the foot in babies or neo-nates.

The present invention overcomes or mitigates the disadvantages of the prior art by using a plurality of closely associated transmitters and generators which allows an "average" evened out signal to be produced and is capable of utilising the non-pulsatile element of the patient's blood flow. In addition, a further advantage of the invention is that it allows the measurement of oxygen saturation (SO_2). Furthermore, the invention permits the measurement of haemoglobin index (HbI) and/or temperature.

According to the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part.

A preferred embodiment of the invention is one wherein the non-pulsatile element of the patient's blood is utilised. Thus, according to a further feature of the invention we provide a device for the non-invasive measurement of one or more analytes in blood in a patient's body part as herein before described which is adapted to measure the non-pulsatile element of a patient's blood. In a further preferred embodiment the

CLAIMS

1. A device for the non-invasive measurement of one or more analytes in blood in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibres positioned to transmit light to the body part and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part.

2. A device according to Claim 1 characterised in that it is adapted to utilise the non-pulsatile element of a patient's blood.

3. A device according to Claim 1 characterised in that it is adapted to utilise the non-pulsatile and the pulsatile elements of a patient's blood.

4. A device according to Claim 2 characterised in that it is so adapted by being provided with a plurality of closely associated transmitters and generators which allow an "average" evened out signal to be produced.

5. A device according to Claim 1 characterised in that it is adapted to measure blood glucose levels.

6. A device according to Claim 1 characterised in that it is adapted to measure blood oxygen saturation (SO_2).

7. A device according to Claim 1 characterised in that it is adapted to measure the haemoglobin index (HbI).

8. A device according to Claim 1 characterised in that it is adapted to measure the temperature of a patient's blood.

9. A device according to Claim 1 characterised in that it is adapted to measure two or more analytes in blood, which analytes are selected from blood glucose levels, blood oxygen saturation (SO_2), the haemoglobin index (HbI) and the temperature of a patient's blood.

10. A device according to Claim 9 characterised in that it is adapted to measure blood glucose levels, blood oxygen saturation (SO_2), the haemoglobin index (HbI) and the temperature of a patient's blood.

11. A device according to Claim 1 characterised in that it is adapted to measure of one or more analytes in blood in a patient's finger or thumb.

12. A device according to Claim 1 characterised in that it is provided with a greater number of transmitter fibres than detector fibres.

13. A device according to Claim 1 characterised in that it is provided with from 12 to 24 transmitter fibres.

14. A device according to Claim 13 characterised in that it is provided with 18 transmitter fibres.

15. A device according to Claim 1 characterised in that it is provided with from 6 to 18 detector fibres.

16. A device according to Claim 14 characterised in that it is provided with 12 detector fibres.

17. A device according to Claim 1 characterised in that diameter of the fibres is from 200 - 300μm.

18. A device according to Claim 1 characterised in that diameter of the fibres is 250μm.

19. A device according to Claim 1 characterised in that the detector fibres are positioned to detect transmitted light rather than reflected light.

20. A device according to Claim 1 characterised in that the wavelength used in the transmitter fibres is from 500 to 1100nm.

21. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is from 800 to 1100nm.

22. A device according to Claim 21 characterised in that the wavelength used in the transmitter fibres is 805nm, 925nm, 970nm and the broadband average 1000-1100nm.

23. A device according to Claim 21 characterised in that the wavelengths are sampled at regular intervals of 1.96nm in the entire range 800nm to 1100nm.

24. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is from 500 to 600nm.

25. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is 500.9nm, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm.

26. A device according to Claim 20 characterised in that the wavelength used in the transmitter fibres is 500.9nm, 528.1nm, 549.5nm, 561.1nm, 572.7nm and 586.3nm; and from 800nm to 1100nm.

27. A method of measuring blood glucose levels which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part.

28. A method according to Claim 27 characterised in that the non-pulsatile element is used.

29. A method according to Claim 28 characterised in that the non-pulsatile and pulsatile element is used.

30. A device according to Claim 1 programmed so as to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

31. A device according to Claim 30 programmed so as to conduct a multiple linear regression analysis on spectral data collected by the detectors.

32. A device according to Claim 30 wherein the Haemoglobin Index is calculated using the equation:

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100$$

where a = absorption value at 500.9nm
b = absorption value at 528.1nm
c = absorption value at 549.5nm
e = absorption value at 572.7nm.

33. A device according to Claim 30 wherein the Oxygenation Index is calculated using the equation:

$$OXI=(ee-d)/11.7-(d-c)/11.6)*100/HbI$$

where c = absorption value at 549.5nm

d = absorption value at 561.1nm

e = absorption value at 572.7nm.

34. A device according to Claim 30 wherein the Oxygen Saturation (SO_2) is calculated using the equation:

$$SO_2=100*(OXI+0.43)/1.5.$$

35. A method of measuring blood oxygen saturation which comprises placing a non-invasive measuring device as herein before described against a body part of a patient and using the detector to measure the light transmitted through or reflected from the body part using a device according to Claim 34.

36. A computer programme product comprising a computer readable medium having thereon computer programme code means, when said programme is loaded, to make the computer execute a procedure to calculate one or more of the haemoglobin index, the oxygen index and the blood oxygen saturation.

37. A computer programme according to Claim 36 wherein the computer programme code means will make the computer execute a procedure to calculate one or more of :

$$HbI = ((b-a)/27.1+(c-b)/21.4+(c-e)/23.3+(c-f)/13.6)*100;$$

$$OXI=(e-d)/11.7-(d-c)/11.6)*100/HbI; \text{ and}$$

$$SO_2=100*(OXI+0.43)/1.5$$

where a = absorption value at 500.9nm

b = absorption value at 528.1nm

c = absorption value at 549.5nm

d = absorption value at 561.1nm

e = absorption value at 572.7nm

38. A device substantially as described with reference to the accompanying examples and drawings.

P36002WO.1

09/743206

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

(PCT Article 36 and Rule 70)

Applicant's or agent's file reference	FOR FURTHER ACTION	See Notification of Transmittal of International Preliminary Examination Report (Form PCT/IPEA/416)
SPG/P36002WO		
International application No.	International filing date *(day/month/year)*	Priority date *(day/month/year)*
PCT/GB99/02127	02/07/1999	04/07/1998
International Patent Classification (IPC) or national classification and IPC A61B5/00		
Applicant WHITLAND RESEARCH LIMITED et al.		

1. This international preliminary examination report has been prepared by this International Preliminary Examining Authority and is transmitted to the applicant according to Article 36.

2. This REPORT consists of a total of 7 sheets, including this cover sheet.

   ☒ This report is also accompanied by ANNEXES, i.e. sheets of the description, claims and/or drawings which have been amended and are the basis for this report and/or sheets containing rectifications made before this Authority (see Rule 70.16 and Section 607 of the Administrative Instructions under the PCT).

   These annexes consist of a total of 7 sheets.

3. This report contains indications relating to the following items:

   - I ☒ Basis of the report
   - II ☐ Priority
   - III ☒ Non-establishment of opinion with regard to novelty, inventive step and industrial applicability
   - IV ☐ Lack of unity of invention
   - V ☒ Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations suporting such statement
   - VI ☐ Certain documents cited
   - VII ☐ Certain defects in the international application
   - VIII ☒ Certain observations on the international application

Date of submission of the demand	Date of completion of this report
01/02/2000	13.10.2000
Name and mailing address of the international preliminary examining authority: European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx: 523656 epmu d Fax: +49 89 2399 - 4465	Authorized officer Fontenay, P Telephone No. +49 89 2399 2646

Form PCT/IPEA/409 (cover sheet) (January 1994)

INTERNATIONAL PRELIMINARY EXAMINATION REPORT

International application No. PCT/GB99/02127

I. Basis of the report

1. This report has been drawn on the basis of (*substitute sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to the report since they do not contain amendments.*):

Description, pages:

1-4,6-12	as originally filed	
5,5a	with telefax of	01/08/2000

Claims, No.:

1-32	with telefax of	01/08/2000

Drawings, sheets:

1/2,2/2	as originally filed

2. The amendments have resulted in the cancellation of:

☐ the description, pages:
☐ the claims, Nos.:
☐ the drawings, sheets:

3. ☐ This report has been established as if (some of) the amendments had not been made, since they have been considered to go beyond the disclosure as filed (Rule 70.2(c)):

4. Additional observations, if necessary:

III. Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:

☐ the entire international application.

☒ claims Nos. 7,32.

because:

☐ the said international application, or the said claims Nos. relate to the following subject matter which does not require an international preliminary examination (*specify*):

☒ the description, claims or drawings (*indicate particular elements below*) or said claims Nos. 7 32 are so unclear that no meaningful opinion could be formed (*specify*):

see separate sheet

☐ the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.

☐ no international search report has been established for the said claims Nos. .

V. Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

1. Statement

Novelty (N)	Yes:	Claims	2-6, 8-22, 27-31
	No:	Claims	1, 23-26
Inventive step (IS)	Yes:	Claims	27-31
	No:	Claims	2-6, 8-22
Industrial applicability (IA)	Yes:	Claims	1-32
	No:	Claims	

2. Citations and explanations

see separate sheet

VIII. Certain observations on the international application

The following observations on the clarity of the claims, description, and drawings or on the question whether the claims are fully supported by the description, are made:

see separate sheet

Re Item III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

The subject-matter of claims 7 and 32 is not clearly defined (Article 6 PCT).

III.1 The wording "characterized in that it is adapted to measure the temperature of a patient's blood" as it appears in claim 7 is ambiguous. It could suggest that the temperature is obtained from the measurements carried out within the claimed device in the same way as it calculates the haemoglobin index or the blood oxygen saturation or that it contains a thermal sensor as it is suggested on page 10, lines 2-4 of the description.

III.2 The claims should not refer to the drawings or to parts of the description (Rule 6.2 (a) PCT). Claim 32 is accordingly not allowable.

Re Item V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement

Reference is made to the following documents:

D1 = WO-A-9727800
D2 = US-A-5755226

V.1 The subject-matter of independent claim 1 is not new in the sense of Article 33(2) PCT.

D2 discloses a device for the non-invasive measurement of one or more analyte in blood (haematocrit) in a patient's body part which comprises a light transmitter comprising a plurality of transmitting fibers positioned to transmit light to the body part (see D2, column 15, line 44 - column 16, line 17, figure 4) and a light detector comprising a plurality of light detector fibres position to detect light transmitted through or reflected from the body part (see D2, column 13, lines 47-62). Moreover, the device disclosed in D2 is adapted to utilise the non-pulsatile element of a patient's blood (see D2, column 6, lines 27-47; column 8, lines 53 -

column 9, line 27). It is here noted that the fact that the claimed device is adapted to utilise the non-pulsatile element of a patient's blood does not exclude that it utilises additional components.

It follows that all the features of claim 1 are known in combination from D2. The subject-matter of claim 1 is accordingly not new.

The device of D2 is programmed so as to calculate a multiplicity of analytes concentration and in particular haematocrit and oxygen saturation. Claim 25 is accordingly also not new. It is also proposed in D1 to base the analysis on multiple linear regression so that claim 26 is also not new.

V.2 The features of dependent claims 2-6, 8-18 and 20-22 are known from or rendered obvious by the prior art as illustrated by documents D1 or D2 and as indicated in the International Search Report in relation with original claims 3-7, 9-12, 15, 17, 19, 21, 22 and 24-26. The dependent claims 2-6, 8-17, and 20-22 therefore do not appear to contain any additional features which, in combination with the features of claim 1 to which they refer, would involve an inventive step.

It is in particular noted that in D2, a plurality of associated transmitters and generators are used and that an average signal is produced so as to consider the non-pulsatile component of the signal (see D2, column 23, line 1- column 24, line 53).

V.3 The feature of the sampling interval of 1,96 nm as it appears in claim 19, constitutes a constructional detail as to the device in order to obtain the information over the whole spectra. This feature cannot justify a positive inventive step assessment since the skilled man will arrive at something falling under the terms of the claims by the exercise of routine trials according to the intended purpose i.e. in order to obtain sufficient information over said spectra. The applicant may also refer to the Guidelines, PCT/GL/3 Chapter IV, 8.8 (C1)(ii).

V.4 The subject-matter of claims 23 and 24, as may be understood (see comments under point VIII) is already known from the prior art. Document D2 discloses a method of measuring blood glucose which comprises placing a non-invasive

measuring device as claimed against a body part (a finger) of a patient and using the detector to measure the light reflected from the body part (see D2, figures 2, 4, 7). In D2, as explained above under section V.1, the non-pulsatile component of the signal is utilised.

The subject-matter of claims 23 and 24 is therefore not new considering the teaching of D2.

V.5 The subject-matter of claim 27 and 31 is not disclosed as such in D2. None of the documents cited in the search report suggest to calculate the Haemoglobin index on the basis of the equation indicated in claim 27 or 31. The subject-matter of claim 27 is accordingly considered to be new and inventive in the sense of article 33(2) and 33(3) PCT considering the prior art as presently known.

Claim 28, when depending on claim 27 (see comments under point VIII.3) would accordingly also be new and inventive. The same would apply to claim 29 when depending on claim 28 (see the comments under point VIII.3).

V.6 The method defined in claim 30 seems to refer to the use of a device as defined in claim 29 (see comments under section VIII). The device of claim 29 being considered as new and inventive, when depending on claim 28, the same applies to said use.

Re Item VIII Certain observations on the international application

VIII.1 The subject-matter of claim 1 is not clearly defined (Article 6 PCT). The wording in the characterising part of claim 1: "is adapted to utilise the non-pulsatile element of a patient's blood is too vague because of the terms "adapted to utilise" which presents a broad meaning. It is in particular considered that any device which senses a signal with a pulsatile and a non-pulsatile component may be considered as being adapted to utilise the non-pulsatile component. The present meaning does not permit to "define" the matter for which protection is sought as requested under Article 6.

VIII.2 The subject-matter of claims 23 and 30 is not clearly defined (Article 6 PCT) because it refers to the device as "herein before described". Said wording is not allowable in the claims. It has been assumed in section V above, that the wording "measuring device according to any of the preceding claims" had been intended.

Claim 30 is also not clear because it refers to a first device (as herein before described) and then to a device according to claim 29.

VIII.3 Claim 28 is not clear because it refers to claim 30. Moreover, claim 28 refers to the parameter HBI which is defined in claim 27. The same applies to claim 29 which refers to the parameter $S0_2$ which is defined in claim 28. It has accordingly been assumed in the present report that claim 28 refers to claim 27 and that claim 29 refers to claim 28.

PATENT COOPERATION TREA

PCT

09/743206

INTERNATIONAL SEARCH REPORT

(PCT Article 18 and Rules 43 and 44)

Applicant's or agent's file reference	FOR FURTHER ACTION	see Notification of Transmittal of International Search Report (Form PCT/ISA/220) as well as, where applicable, item 5 below.
SPG/P36002WO		
International application No.	International filing date *(day/month/year)*	(Earliest) Priority Date *(day/month/year)*
PCT/GB 99/02127	02/07/1999	04/07/1998

Applicant

WHITLAND RESEARCH LIMITED et al.

This International Search Report has been prepared by this International Searching Authority and is transmitted to the applicant according to Article 18. A copy is being transmitted to the International Bureau.

This International Search Report consists of a total of 3 sheets.

[X] It is also accompanied by a copy of each prior art document cited in this report.

1. **Basis of the report**

   a. With regard to the **language**, the international search was carried out on the basis of the international application in the language in which it was filed, unless otherwise indicated under this item.

   [ ] the international search was carried out on the basis of a translation of the international application furnished to this Authority (Rule 23.1(b)).

   b. With regard to any **nucleotide and/or amino acid sequence** disclosed in the international application, the international search was carried out on the basis of the sequence listing :

   [ ] contained in the international application in written form.

   [ ] filed together with the international application in computer readable form.

   [ ] furnished subsequently to this Authority in written form.

   [ ] furnished subsequently to this Authority in computer readble form.

   [ ] the statement that the subsequently furnished written sequence listing does not go beyond the disclosure in the international application as filed has been furnished.

   [ ] the statement that the information recorded in computer readable form is identical to the written sequence listing has been furnished

2. [X] **Certain claims were found unsearchable** (See Box I).

3. [ ] **Unity of invention is lacking** (see Box II).

4. With regard to the **title,**

   [ ] the text is approved as submitted by the applicant.

   [X] the text has been established by this Authority to read as follows:

   NON-INVASIVE MEASUREMENT OF BLOOD ANALYTES

5. With regard to the **abstract,**

   [X] the text is approved as submitted by the applicant.

   [ ] the text has been established, according to Rule 38.2(b), by this Authority as it appears in Box III. The applicant may, within one month from the date of mailing of this international search report, submit comments to this Authority.

6. The figure of the **drawings** to be published with the abstract is Figure No. _____

   [ ] as suggested by the applicant.

   [ ] because the applicant failed to suggest a figure.

   [ ] because this figure better characterizes the invention.

   [X] None of the figures.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB 99/02127

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. [X] Claims Nos.: 36,37
because they relate to subject matter not required to be searched by this Authority, namely:
Rule 39.1(iv) PCT - Program for computers

2. [ ] Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:

3. [ ] Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. [ ] As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. [ ] As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. [ ] As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. [ ] No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

[ ] The additional search fees were accompanied by the applicant's protest.

[ ] No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/GB 99/02127

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61B5/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 5 755 226 A (3M) 26 May 1998 (1998-05-26) the whole document ---	1-4,6,7, 9-11,15, 19-22, 24-31,38
X	WO 97 27800 A (DIASENSE) 7 August 1997 (1997-08-07) the whole document -----	1,5,12, 15-17

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

° Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
27 September 1999	05/10/1999

Name and mailing address of the ISA	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Lemercier, D

1

INTERNATIONAL SEARCH REPORT

[redacted]mation on patent family members

[redacted]ternational Application No
PCT/GB 99/02127

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 5755226 A	26-05-1998	US 5553615 A	10-09-1996
		EP 0742896 A	20-11-1996
		JP 9508291 T	26-08-1997
		WO 9520757 A	03-08-1995
WO 9727800 A	07-08-1997	AU 1846897 A	22-08-1997
		EP 0889703 A	13-01-1999